following concurrent resolution, which was referred to the Committee on Health, Education, Labor, and Pensions:

#### S. CON. RES. 33

Whereas school music programs enhance intellectual development and enrich the academic environment for students of all ages;

Whereas students who participate in school music programs are less likely to be involved with drugs, gangs, or alcohol, and have better attendance in school;

Whereas the skills gained through sequential music instruction, including discipline and the ability to analyze, solve problems, communicate, and work cooperatively, are vital for success in the 21st century work-place:

Whereas the majority of students attending public schools in inner city neighborhoods have virtually no access to music education, which places them at a disadvantage compared to their peers in other communities;

Whereas the arts are a core academic subject, and music is an essential element of the arts; and

Whereas every student in the United States should have an opportunity to reap the benefits of music education: Now, therefore, be it

Resolved by the Senate (the House of Representatives concurring), That it is the sense of Congress that music education grounded in rigorous instruction is an important component of a well-rounded academic curriculum and should be available to every student in every school in the United States.

# $\begin{array}{c} {\rm AMENDMENTS~SUBMITTED~AND} \\ {\rm PROPOSED} \end{array}$

SA 1045. Mr. REID (for Mr. OBAMA) submitted an amendment intended to be proposed by Mr. REID to the bill S. 1082, to amend the Federal Food, Drug, and Cosmetion Act to reauthorize and amend the prescription drug user fee provisions, and for other purposes; which was ordered to lie on the table.

SA 1046. Ms. STABENOW (for herself, Mr. KOHL, Mr. HATCH, and Mr. COBURN) submitted an amendment intended to be proposed by her to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1047. Mr. ROBERTS (for himself, Mr. HARKIN, Mr. BURR, and Mr. COBURN) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1048. Ms. SNOWE submitted an amendment intended to be proposed by her to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1049. Mr. ENZI (for himself and Mr. Kennedy) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

\$A 1050. Mr. ENZI (for himself and Mr. Kennedy) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1051. Mr. STEVENS (for himself and Ms. Murkowski) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the

SA 1052. Mr. CORKER submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1053. Mr. ENZI (for himself, Mr. Kennedy, Mr. Dodd, and Mrs. CLINTON) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1054. Mr. FEINGOLD submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1055. Mr. LEVIN submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1056. Mr. REED (for himself and Mr. ISAKSON) submitted an amendment intended to be proposed by him to the bill S. 1082, supra: which was ordered to lie on the table.

\$A 1057. Mr. GREGG submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table

SA 1058. Mr. DEMINT (for himself, Mr. COBURN, and Mr. MARTINEZ) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1059. Mr. SESSIONS (for himself, Mrs. LINCOLN, Mr. COCHRAN, Mr. PRYOR, Mr. LOTT, and Mr. SHELBY) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

SA 1060. Mr. HATCH (for himself and Mr. Kennedy) submitted an amendment intended to be proposed by him to the bill S. 1082, supra; which was ordered to lie on the table.

### TEXT OF AMENDMENTS

SA 1045. Mr. REID (for Mr. OBAMA) submitted an amendment intended to be proposed by Mr. Reid to the bill S. 1082, to amend the Federal Food, Drug, and Cosmetic Act to reauthorize and amend the prescription drug user fee provisions, and for other purposes; which was ordered to lie on the table; as follows:

At the appropriate place, insert the following:

### SEC. \_\_\_. IMPROVING GENETIC TEST SAFETY AND QUALITY.

Not later than 30 days after the date of enactment of this Act, the Secretary shall enter into a contract with the Institute of Medicine to conduct a study to assess the overall safety and quality of genetic tests and prepare a report that includes recommendations to improve Federal oversight and regulation of genetic tests. Such study shall take into consideration relevant reports by the Secretary's Advisory Committee on Genetic Testing and other groups and shall be completed not later than 1 year after the date on which the Secretary entered into such contract.

SA 1046. Ms. STABENOW (for herself, Mr. Kohl, Mr. Hatch, and Mr. Coburn) submitted an amendment intended to be proposed by her to the bill S. 1082, to amend the Federal Food, Drug, and Cosmetic Act to reauthorize and amend the prescription drug user fee provisions, and for other purposes; which was ordered to lie on the table; as follows:

At the appropriate place, insert the following:

## SEC. \_\_\_. CITIZENS PETITIONS AND PETITIONS FOR STAY OF AGENCY ACTION.

Section 505 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355), as amended by this Act, is amended by adding at the end the following:

"(s) CITIZEN PETITIONS AND PETITIONS FOR STAY OF AGENCY ACTION.—

"(1) IN GENERAL.—

``(A) No delay of consideration or approval.—

"(i) IN GENERAL.—With respect to a pending application submitted under subsection (b)(2) or (j), if a petition is submitted to the Secretary that seeks to have the Secretary take, or refrain from taking, any form of action relating to the approval of the application, including a delay in the effective date of the application, clauses (ii) and (iii) shall apply.

"(ii) NO DELAY OF CONSIDERATION OR AP-PROVAL.—Except as provided in clause (iii), the receipt and consideration of a petition described in clause (i) shall not delay consideration or approval of an application submitted under subsection (b)(2) or (j).

"(iii) NO DELAY OF APPROVAL WITHOUT DETERMINATION.—The Secretary shall not delay approval of an application submitted under subsection (b)(2) or (j) while a petition described in clause (i) is reviewed and considered unless the Secretary determines, not later than 25 business days after the submission of the petition, that a delay is necessary to protect the public health.

"(B) DETERMINATION OF DELAY.—With respect to a determination by the Secretary under subparagraph (A)(iii) that a delay is necessary to protect the public health the following shall apply:

"(i) Not later than 5 days after making such determination, the Secretary shall publish on the Internet website of the Food and Drug Administration a detailed statement providing the reasons underlying the determination. The detailed statement shall include a summary of the petition and comments and supplements, the specific substantive issues that the petition raises which need to be considered prior to approving a pending application submitted under subsection (b)(2) or (j), and any clarifications and additional data that is needed by the Secretary to promptly review the petition.

"(ii) Not later than 10 days after making such determination, the Secretary shall provide notice to the sponsor of the pending application submitted under subsection (b)(2) or (j) and provide an opportunity for a meeting with appropriate staff as determined by the Commissioner to discuss the determination.

''(2) TIMING OF FINAL AGENCY ACTION ON PETITIONS.—

"(A) IN GENERAL.—Notwithstanding a determination made by the Secretary under paragraph (1)(A)(iii), the Secretary shall take final agency action with respect to a petition not later than 180 days of submission of that petition unless the Secretary determines, prior to the date that is 180 days after the date of submission of the petition, that a delay is necessary to protect the public health.

"(B) DETERMINATION OF DELAY.—With respect to a determination by the Secretary under subparagraph (A) that a delay is necessary to protect the public health the following shall apply:

"(i) Not later than 5 days after making the determination under subparagraph (A), the Secretary shall publish on the Internet website of the Food and Drug Administration a detailed statement providing the reasons underlying the determination. The detailed statement should include the state of the review of the petition, the specific outstanding issues that still need to be resolved, a proposed timeframe to resolve the issues, and any additional information that has been requested by the Secretary of the petitioner or needed by the Secretary in order to resolve the petition and not further delay an application filed under subsection (b)(2) or (j).

"(ii) Not later than 10 days after making the determination under subparagraph (A), the Secretary shall provide notice to the sponsor of the pending application submitted